



RAW SEQUENCE LISTING ERROR REPORT

1-3
BIOTECHNOLOGY
SYSTEMS
BRANCH



~~0230~~
#4

The Biotechnology Systems Branch of the Scientific and Technical Information Center (STIC) detected errors when processing the following computer readable form:

Application Serial Number: 09/659,737

Source: OIPE

Date Processed by STIC: 2-16-01

RECEIVED
FEB 28 2001
OIPE/JCWS

THE ATTACHED PRINTOUT EXPLAINS DETECTED ERRORS.

PLEASE FORWARD THIS INFORMATION TO THE APPLICANT BY EITHER:

- 1) INCLUDING A COPY OF THIS PRINTOUT IN YOUR NEXT COMMUNICATION TO THE APPLICANT, WITH A NOTICE TO COMPLY or,
- 2) TELEPHONING APPLICANT AND FAXING A COPY OF THIS PRINTOUT, WITH A NOTICE TO COMPLY

FOR CRF SUBMISSION QUESTIONS, PLEASE CONTACT MARK SPENCER, 703-308-4212.

FOR SEQUENCE RULES INTERPRETATION, PLEASE CONTACT ROBERT WAX, 703-308-4216.

PATENTIN 2.1 e-mail help: patin21help@uspto.gov or phone 703-306-4119 (R. Wax)

PATENTIN 3.0 e-mail help: patin3help@uspto.gov or phone 703-306-4119 (R. Wax)

TO REDUCE ERRORED SEQUENCE LISTINGS, PLEASE USE THE CHECKER
VERSION 3.0 PROGRAM, ACCESSIBLE THROUGH THE U.S. PATENT AND
TRADEMARK OFFICE WEBSITE. SEE BELOW:

Checker Version 3.0

The Checker Version 3.0 application is a state-of-the-art Windows based software program employing a logical and intuitive user-interface to check whether a sequence listing is in compliance with format and content rules. Checker Version 3.0 works for sequence listings generated for the original version of 37 CFR §§1.821 - 1.825 effective October 1, 1990 (old rules) and the revised version (new rules) effective July 1, 1998 as well as World Intellectual Property Organization (WIPO) Standard ST 25.

Checker Version 3.0 replaces the previous DOS-based version of Checker, and is Y2K-compliant. Checker allows public users to check sequence listings in Computer Readable form (CRF) before submitting them to the United States Patent and Trademark Office (USPTO). Use of Checker prior to filing the sequence listing is expected to result in fewer errored sequence listings, thus saving time and money.

Checker Version 3.0 can be down loaded from the USPTO website at the following address:
<http://www.uspto.gov/web/offices/pac/checker>

Sequence Listing Error Summary

ERROR DETECTED SUGGESTED CORRECTION

SERIAL NUMBER: 09/659,737

ATTN: NEW RULES CASES: PLEASE DISREGARD ENGLISH "ALPHA" HEADERS, WHICH WERE INSERTED BY PTO SOFTWARE

1 ☐ Wrapped Nucleics

The number/text at the end of each line "wrapped" down to the next line.
This may occur if your file was retrieved in a word processor after creating it.
Please adjust your right margin to .3, as this will prevent "wrapping".

2 ☐ Wrapped Aminos

The amino acid number/text at the end of each line "wrapped" down to the next line.
This may occur if your file was retrieved in a word processor after creating it.
Please adjust your right margin to .3, as this will prevent "wrapping".

3 ☐ Incorrect Line Length

The rules require that a line not exceed 72 characters in length. This includes spaces.

4 ☐ Misaligned Amino Acid Numbering

The numbering under each 5th amino acid is misaligned. This may be caused by the use of tabs between the numbering. It is recommended to delete any tabs and use spacing between the numbers.

5 ☐ Non-ASCII

This file was not saved in ASCII (DOS) text, as required by the Sequence Rules.
Please ensure your subsequent submission is saved in ASCII text so that it can be processed.

6 ☐ Variable Length

Sequence(s) ☐ contain n's or Xaa's which represented more than one residue.
As per the rules, each n or Xaa can only represent a single residue.
Please present the maximum number of each residue having variable length and indicate in the (ix) feature section that some may be missing.

7 ☐ PatentIn ver. 2.0 "bug"

A "bug" in PatentIn version 2.0 has caused the <220>-<223> section to be missing from amino acid sequence(s) ☐. Normally, PatentIn would automatically generate this section from the previously coded nucleic acid sequence. Please manually copy the relevant <220>-<223> section to the subsequent amino acid sequence. This applies primarily to the mandatory <220>-<223> sections for Artificial or Unknown sequences.

8 ☐ Skipped Sequences (OLD RULES)

Sequence(s) ☐ missing. If intentional, please use the following format for each skipped sequence:
(2) INFORMATION FOR SEQ ID NO:X:
(i) SEQUENCE CHARACTERISTICS:(Do not insert any headings under "SEQUENCE CHARACTERISTICS")
(xi) SEQUENCE DESCRIPTION:SEQ ID NO:X:
This sequence is intentionally skipped

Please also adjust the "(iii) NUMBER OF SEQUENCES:" response to include the skipped sequence(s).

9 ☐ Skipped Sequences (NEW RULES)

Sequence(s) ☐ missing. If intentional, please use the following format for each skipped sequence.
<210> sequence id number
<400> sequence id number
000

10 ☒ Use of n's or Xaa's (NEW RULES)

Use of n's and/or Xaa's have been detected in the Sequence Listing.
Use of <220> to <223> is MANDATORY if n's or Xaa's are present.
In <220> to <223> section, please explain location of n or Xaa, and which residue n or Xaa represents.

11 ☐ Use of <213>Organism (NEW RULES)

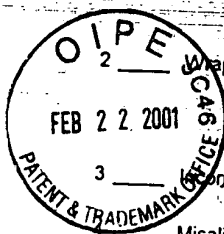
Sequence(s) ☐ are missing this mandatory field or its response.

12 ☐ Use of <220>Feature (NEW RULES)

Sequence(s) ☐ are missing the <220>Feature and associated headings.
Use of <220> to <223> is MANDATORY if <213>ORGANISM is "Artificial" or "Unknown"
Please explain source of genetic material in <220> to <223> section.
(See "Federal Register," 6/01/98, Vol. 63, No. 104, pp. 29631-32) (Sec. 1.823 of new Rules)

13 ☐ PatentIn ver. 2.0 "bug"

Please do not use "Copy to Disk" function of PatentIn version 2.0. This causes a corrupted file, resulting in missing mandatory numeric identifiers and responses (as indicated on raw sequence listing). Instead, please use "File Manager" or any other means to copy file to floppy disk.





OIPE

RAW SEQUENCE LISTING
 PATENT APPLICATION: US/09/659,737

DATE: 02/16/2001
 TIME: 11:33:38

Input Set : A:\10489a1.app
 Output Set: N:\CRF3\02162001\I659737.raw

Does Not Comply
 Corrected Diskette Needed
 see p. 6

3 <110> APPLICANT: Blumenberg, Miroslav
 4 Gazel, Alix M
 6 <120> TITLE OF INVENTION: GENES AND POLYNUCLEOTIDES ASSOCIATED WITH ULTRAVIOLET
 7 RADIATION-MEDIATED SKIN DAMAGE AND USES THEREOF
 9 <130> FILE REFERENCE: PC10489A
 C--> 11 <140> CURRENT APPLICATION NUMBER: US/09/659,737
 C--> 12 <141> CURRENT FILING DATE: 2000-09-11
 14 <150> PRIOR APPLICATION NUMBER: 60/155,029
 15 <151> PRIOR FILING DATE: 1999-09-20
 17 <160> NUMBER OF SEQ ID NOS: 19
 19 <170> SOFTWARE: PatentIn Ver. 2.1
 21 <210> SEQ ID NO: 1
 22 <211> LENGTH: 164
 23 <212> TYPE: DNA
 24 <213> ORGANISM: Homo sapiens
 26 <220> FEATURE:
 27 <221> NAME/KEY: CDS
 28 <222> LOCATION: (2)..(163)
 30 <400> SEQUENCE: 1
 31 g cac cgg gac atc aag gca gga aat att ttg cta ctt gag aag ata gaa 49
 32 His Arg Asp Ile Lys Ala Gly Asn Ile Leu Leu Leu Glu Lys Ile Glu
 33 1 5 10 15
 35 cat gat gac atc tgc aat aaa act ttg aag att aca gat ttt ggg ttg 97
 36 His Asp Asp Ile Cys Asn Lys Thr Leu Lys Ile Thr Asp Phe Gly Leu
 37 20 25 30
 39 gcg agg gaa tgg cac agg acc acc aaa atg agc aca gca ggc acc tat 145
 40 Ala Arg Glu Trp His Arg Thr Thr Lys Met Ser Thr Ala Gly Thr Tyr
 41 35 40 45
 43 gcc tgg atg gcc cca gaa'g
 44 Ala Trp Met Ala Pro Glu 164
 45 50
 48 <210> SEQ ID NO: 2
 49 <211> LENGTH: 54
 50 <212> TYPE: PRT
 51 <213> ORGANISM: Homo sapiens
 53 <400> SEQUENCE: 2
 54 His Arg Asp Ile Lys Ala Gly Asn Ile Leu Leu Leu Glu Lys Ile Glu
 55 1 5 10 15
 57 His Asp Asp Ile Cys Asn Lys Thr Leu Lys Ile Thr Asp Phe Gly Leu
 58 20 25 30
 60 Ala Arg Glu Trp His Arg Thr Thr Lys Met Ser Thr Ala Gly Thr Tyr
 61 35 40 45
 63 Ala Trp Met Ala Pro Glu
 64 50
 68 <210> SEQ ID NO: 3
 69 <211> LENGTH: 145
 70 <212> TYPE: DNA

RAW SEQUENCE LISTING

PATENT APPLICATION: US/09/659,737

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TIME: 11:33:38

Input Set : A:\10489a1.app

Output Set: N:\CRF3\02162001\I659737.raw

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71 <213> ORGANISM: Homo sapiens
73 <220> FEATURE:
74 <221> NAME/KEY: CDS
75 <222> LOCATION: (2)..(145)
77 <400> SEQUENCE: 3
78 a\cat cgg gac atc aag agc gac tcg atc ctg ctg acc cat gat ggc agg 49
79 His Arg Asp Ile Lys Ser Asp Ser Ile Leu Leu Thr His Asp Gly Arg
80 1 5 10 15
82 gtg aag ctg tca gac ttt ggg ttc tgc gcc cag gtg agc aag gaa gtg 97
83 Val Lys Leu Ser Asp Phe Gly Phe Cys Ala Gln Val Ser Lys Glu Val
84 20 25 30
86 ccc cga agg aag tcg ctg gtc ggc acg ccc tac tgg atg gcc cca gag 145
87 Pro Arg Arg Lys Ser Leu Val Gly Thr Pro Tyr Trp Met Ala Pro Glu
88 35 40 45
91 <210> SEQ ID NO: 4
92 <211> LENGTH: 48
93 <212> TYPE: PRT
94 <213> ORGANISM: Homo sapiens
96 <400> SEQUENCE: 4
97 His Arg Asp Ile Lys Ser Asp Ser Ile Leu Leu Thr His Asp Gly Arg
98 1 5 10 15
100 Val Lys Leu Ser Asp Phe Gly Phe Cys Ala Gln Val Ser Lys Glu Val
101 20 25 30
103 Pro Arg Arg Lys Ser Leu Val Gly Thr Pro Tyr Trp Met Ala Pro Glu
104 35 40 45
108 <210> SEQ ID NO: 5
109 <211> LENGTH: 146
110 <212> TYPE: DNA
111 <213> ORGANISM: Homo sapiens
113 <220> FEATURE:
114 <221> NAME/KEY: CDS
115 <222> LOCATION: (2)..(145)
117 <400> SEQUENCE: 5
118 t\cac agg gac atc aag agt gac tcc atc ctg ctg acc ctc gat ggc agg 49
119 His Arg Asp Ile Lys Ser Asp Ser Ile Leu Leu Thr Leu Asp Gly Arg
120 1 5 10 15
122 gtg aag ctc tcg gac ttc gga ttc tgt gct cag atc agc aaa gac gtc 97
123 Val Lys Leu Ser Asp Phe Gly Phe Cys Ala Gln Ile Ser Lys Asp Val
124 20 25 30
126 cct aag agg aag tcc ctg gtg gga acc ccc tac tgg atg gcg ccc gag 146
127 Pro Lys Arg Lys Ser Leu Val Gly Thr Pro Tyr Trp Met Ala Pro Glu
128 35 40 45
131 <210> SEQ ID NO: 6
132 <211> LENGTH: 48
133 <212> TYPE: PRT
134 <213> ORGANISM: Homo sapiens
136 <400> SEQUENCE: 6
137 His Arg Asp Ile Lys Ser Asp Ser Ile Leu Leu Thr Leu Asp Gly Arg
138 1 5 10 15

```

RAW SEQUENCE LISTING

PATENT APPLICATION: US/09/659,737

DATE: 02/16/2001

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Input Set : A:\10489a1.app

Output Set: N:\CRF3\02162001\I659737.raw

```

140 Val Lys Leu Ser Asp Phe Gly Phe Cys Ala Gln Ile Ser Lys Asp Val
141          20          25          30
143 Pro Lys Arg Lys Ser Leu Val Gly Thr Pro Tyr Trp Met Ala Pro Glu
144          35          40          45
148 <210> SEQ ID NO: 7
149 <211> LENGTH: 3627
150 <212> TYPE: DNA
151 <213> ORGANISM: Homo sapiens
153 <220> FEATURE:
154 <221> NAME/KEY: CDS
155 <222> LOCATION: (868)..(1275)
157 <220> FEATURE:
158 <221> NAME/KEY: CDS
159 <222> LOCATION: (1420)..(1553)
161 <220> FEATURE:
162 <221> NAME/KEY: CDS
163 <222> LOCATION: (1900)..(2026)
165 <220> FEATURE:
166 <221> NAME/KEY: CDS
167 <222> LOCATION: (2105)..(2230)
169 <220> FEATURE:
170 <221> NAME/KEY: CDS
171 <222> LOCATION: (2696)..(2833)
173 <400> SEQUENCE: 7
174 gatctgcgac ctccttcaga acctgccaaa atgactagga aaaatgctgt ttccatagca 60
176 agagccaaaa gagaacatga cggccctgca ctccgggato tctctggcac cagattccca 120
178 gccagggga gacacctgaa cccccagat ggtgacacac ctctgtggtc ctctgtcagg 180
180 gacataacct cccagcacag atttgcaaac tccctgctgc aggcacaagc agggctatcg 240
182 gccccaggt gtggtcccc tgccttggtt cagggagtgg agacacagtt gccactgct 300
184 cccacccca ctgccaggcc tcttctgccc ccatgggtcc tggggtgggg gagccttggg 360
186 agttgaagaa tgcctctgac ccagattctt caagcagcct ctgagctcag aggaagagtc 420
188 tgcctcacgg cagcctccct ggggtctagc tgtcaatcgc ccaggaagaa ataccagcg 480
190 cgggaccggg cggggaagct ggccttctct gtcttcccag gtgcagcaca gcgagtgtaa 540
192 ggagctgtct tgggcttgcc cagcctgggtg cctgcgggg gactgctggc acaggactgt 600
194 gactgggctt cagctctgtc tgaaaatctt tgcttcagag cactcccta gtttgatctg 660
196 ataccggcc tgacctgcc agagtccaga ggtcacggcg gccagccct gcctccggga 720
198 aggttatccc aaatgctccc acagccctga ccttctgtg tgetttgtcc cttgcagccc 780
200 aactcctctt tccgaccgcc gcagaaagac aaccccccaa gcctggtggc caaggccag 840
202 tcttgccct cggaccagcc ggtgggg acc ttc agc cct ctg acc act tcy gat 894
203
204          1          5
206 acc agc agc ccc cag aag tcc ctg cgc aca gcc cgg gcc aca ggc cag 942
207 Thr Ser Ser Pro Gln Lys Ser Leu Arg Thr Ala Pro Ala Thr Gly Gln
208 10          15          20          25
210 ctt cca ggc cgg tct tcc cca gcg gga tcc ccc cgc acc tgg cac gcc 990
211 Leu Pro Gly Arg Ser Ser Pro Ala Gly Ser Pro Arg Thr Trp His Ala
212          30          35          40
214 cag atc agc acc agc aac ctg tac ctg ccc cag gac ccc acg gtt gcc 1038
215 Gln Ile Ser Thr Ser Asn Leu Tyr Leu Pro Gln Asp Pro Thr Val Ala

```

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TIME: 11:33:38

Input Set : A:\10489a1.app

Output Set: N:\CRF3\02162001\I659737.raw

```

216          45          50          55
218 aag ggt gcc ctg gct ggt gag gac aca ggt gtt gtg aca cat gag cag 1086
219 Lys Gly Ala Leu Ala Gly Glu Asp Thr Gly Val Val Thr His Glu Gln
220          60          65          70
222 ttc aag gct gcg ctc agg atg gtg gtg gac cag ggt gac ccc cgg ctg 1134
223 Phe Lys Ala Ala Leu Arg Met Val Val Asp Gln Gly Asp Pro Arg Leu
224          75          80          85
226 ctg ctg gac agc tac gtg aag att ggc gag ggc tcc acc ggc atc gtc 1182
227 Leu Leu Asp Ser Tyr Val Lys Ile Gly Glu Gly Ser Thr Gly Ile Val
228 90          95          100          105
230 tgc ttg gcc cgg gaa cac tcg ggc cgc cag gtg gcc gtc aag atg 1230
231 Cys Leu Ala Arg Glu Glu His Ser Gly Arg Gln Val Ala Val Lys Met
232          110          115          120
234 atg gac ctc aga aag cag cag cgc agg gag ctg ctc ttc aac gag / 1275
235 Met Asp Leu Arg Lys Gln Gln Arg Arg Glu Leu Leu Phe Asn Glu
236          125          130          135
238 gtggaggac aggggtgggac acacacgggg gcgttgggga tgggcagtga gcagccagcc 1335
240 aggctggaca tctgtgagca ggggcagtgg gtggccatgc gtctggggcac tgtgcctggc 1395
242 actcaggccc ccacctgccc ccag gtg gtg atc atg cgg gac tac cag cac 1446
243          Val Val Ile Met Arg Asp Tyr Gln His
244          140          145
246 ttc aac gtg gtg gag atg tac aag agc tac ctg gtg ggc gag gag ctg 1494
247 Phe Asn Val Val Glu Met Tyr Lys Ser Tyr Leu Val Gly Glu Glu Leu
248          150          155          160
250 tgg gtg ctc atg gag ttc ctg cag gga gga gcc ctc aca gac atc gtc 1542
251 Trp Val Leu Met Glu Phe Leu Gln Gly Gly Ala Leu Thr Asp Ile Val
252          165          170          175
254 tcc caa gtc ag gtgggcagct gggagggtg gaccctgagt gcaggctgcc 1593
255 Ser Gln Val Arg
256          180
258 ctcaccatgg ccctgccagg gcaatgtggt cttctgcctg tgcccagaa gacttgggat 1653
260 gcctgggctc ccctgcctgc tggggttaact gagaccagg ggtcttgga gtggagaaga 1713
262 gaaggatagc ttctagccaa agctcaggcc ccagttttca ccagggtat ggccgtgactg 1773
264 tgctgcaaaa cagattgcct gggagctgtg gggcctagca ccagggactc ctactctgct 1833
266 cagccacccc acgacctgcc agagctaacg ttctctttca tgggttgccc ccaccttcct 1893
268 gtccag g ctg aat gag gag cag att gcc act gtg tgt gag gct gtg ctg 1942
269          Leu Asn Glu Glu Gln Ile Ala Thr Val Cys Glu Ala Val Leu
270          185          190          195
272 cag gcc ctg gcc tac ctg cat gct cag ggt gtc atc cac cgg gac atc 1990
273 Gln Ala Leu Ala Tyr Leu His Ala Gln Gly Val Ile His Arg Asp Ile
274          200          205          210
276 aag agt gac tcc atc ctg ctg acc ctc gat ggc agg gtaggtccca 2036
277 Lys Ser Asp Ser Ile Leu Leu Thr Leu Asp Gly Arg
278          215          220
280 tctgtccct ggcacagcca cgtcccaact tctcctgat ccaccactca ctcccttttc 2096
282 aaccgcag gtg aag ctc tcg gac ttc gga ttc tgt gct cag atc agc aaa 2146
283          Val Lys Leu Ser Asp Phe Gly Phe Cys Ala Gln Ile Ser Lys
284          225          230          235
286 gac gtc cct aag agg aag tcc ctg gtg gga acc ccc tac tgg atg gct 2194

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RAW SEQUENCE LISTING

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DATE: 02/16/2001

TIME: 11:33:38

Input Set : A:\10489a1.app

Output Set: N:\CRF3\02162001\I659737.raw

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287 Asp Val Pro Lys Arg Lys Ser Leu Val Gly Thr Pro Tyr Trp Met Ala
288      240      245      250
290 cct gaa gtg atc tcc agg tct ttg tat gcc act gag gtaaccgttc 2240
291 Pro Glu Val Ile Ser Arg Ser Leu Tyr Ala Thr Glu
292      255      260      265
294 cctccacccc ccagacctcc caaaagcaac ttggcaactg gcagctcttc tgcgtgtggc 2300
296 cctccagtga gctcaccaaa agcagocctg gttttcagag tcccacctag tcaacaccct 2360
298 tcccccttcc gatggggctg ctcttaccga gtgactttgc tgcagggaac gagtctctga 2420
300 agtgccttcc tcagctcaag ggcagaatgg ggtatggccg ggctctctat gtatgatggc 2480
302 ctctctctga gtgactgaca gctgtgtccc tataggcagt ggtcactcat gcaggcagta 2540
304 actggccaca gggcagggtga ccaggggagg aaggagacag accaccaag gagagctggg 2600
306 gccagctgtc cccctccac cactgctgcc accagaacgc agctaccaat gggccagggt 2660
308 ctggccatgg ggtcaggggac attttctctc tgcag'gtg gat atc tgg tct ctg 2713
309      Val Asp Ile Trp Ser Leu
310      270
312 ggg atc atg gtg att gag atg gta gat ggg gag cca ccg tac ttc agt 2761
313 Gly Ile Met Val Ile Glu Met Val Asp Gly Glu Pro Pro Tyr Phe Ser
314      275      280      285
316 gac tcc cca gtg caa gcc atg aag agg'ctc cgg gac agc ccc cca ccc 2809
317 Asp Ser Pro Val Gln Ala Met Lys Arg Leu Arg Asp Ser Pro Pro Pro
318      290      295      300
320 aag ctg aaa aac tct cac aag gtc'agttggcaca caagggtgcg acctcgcaga 2863
321 Lys Leu Lys Asn Ser His Lys Val
322      305      310
324 cccatttct cctgaggcaa ggggaccaga acctgggctc ccagcatctc ccttccactg 2923
326 aagccacagg gtctgggctc ctggaaaagg ctctcttctc cccacacaaa acccgcacct 2983
328 ggtgtgtggag ccgcatctac gcacaagtgc gcatgtgcgc tccgacaagt cgcctccac 3043
330 ggtgtgtggca ggagagttgc tgcttggcag aagggttgct gcttggcagg cactggtcgg 3103
332 aagcccagtg gggcccata gagggaagg ccaggacacc agcaactccc tgcgttccag 3163
334 ggagggatcc ggagaagctt cactgagcac aaacccttca acccgtgtcg ggagatccat 3223
336 accatgatto gatgtccctg tccatcacgg cgagtcggct catgtcccat tegtgcaca 3283
338 ccccgacaca gctaagccac agcgttcccc ttaaagccag tataagtga tggaagtgg 3343
340 atacatgtaa cctttttgc caaatcgccc ccaacccgc aggccttact gtggacgcc 3403
342 cctgctggca ggtcagcacg gggctgataa gtggcaccgc catctggtgg ccaaaacaag 3463
344 aaatgtctca gagggctgaa gcctctctc taaaatagca aaaaaacaag agttctgtgg 3523
346 ccccaacaca aagctggatg ggaggaccaa caggaaacat ctccaagac aactggtcct 3583
348 tggagcccgc accgctaacc caaaattag catataaagc atgc 3627
351 <210> SEQ ID NO: 8
352 <211> LENGTH: 311
353 <212> TYPE: PRT
354 <213> ORGANISM: Homo sapiens
356 <400> SEQUENCE: 8
357 Thr Phe Ser Pro Leu Thr Thr Ser Asp Thr Ser Ser Pro Gln Lys Ser
358      1      5      10      15
360 Leu Arg Thr Ala Pro Ala Thr Gly Gln Leu Pro Gly Arg Ser Ser Pro
361      20      25      30
363 Ala Gly Ser Pro Arg Thr Trp His Ala Gln Ile Ser Thr Ser Asn Leu
364      35      40      45
366 Tyr Leu Pro Gln Asp Pro Thr Val Ala Lys Gly Ala Leu Ala Gly Glu

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<210> 14
 <211> 18
 <212> DNA
 <213> Homo sapiens

<400> 14
 atgcamcang ayathaar

18

<210> 15
 <211> 20
 <212> DNA
 <213> Homo sapiens

<400> 15
 gcnacytong ngccatcca

<210> 16
 <211> 27
 <212> DNA
 <213> Homo sapiens

<400> 16
 cccgaattca tgcamcanga yathaar

<210> 17
 <211> 29
 <212> DNA
 <213> Homo sapiens

<400> 17
 cccgaattcg cnacytongg ngccatcca

Missing mandatory <220> to <223>
 features to explain²⁰ the "n's" in
 the sequences. See #10 on the
 Error Summary Sheet.

Note: Though not shown²⁷, seq #11
 has an "n" at position 453,
 which needs a <220> to <223>
 explanation.

29

VERIFICATION SUMMARY

PATENT APPLICATION: US/09/659,737

DATE: 02/16/2001

TIME: 11:33:39

Input Set : A:\10489a1.app

Output Set: N:\CRF3\02162001\I659737.raw

L:11 M:270 C: Current Application Number differs, Replaced Application Number
L:12 M:271 C: Current Filing Date differs, Replaced Current Filing Date
L:732 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:11
L:732 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:11
L:732 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:11
L:732 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:11
L:732 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:11
L:1091 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:14
L:1091 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:14
L:1091 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:14
L:1091 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:14
L:1091 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:14
L:1100 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:15
L:1100 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:15
L:1100 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:15
L:1100 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:15
L:1100 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:15
L:1109 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:16
L:1109 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:16
L:1109 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:16
L:1109 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:16
L:1109 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:16
L:1118 M:258 W: Mandatory Feature missing, <220> not found for SEQ ID#:17
L:1118 M:258 W: Mandatory Feature missing, <221> not found for SEQ ID#:17
L:1118 M:258 W: Mandatory Feature missing, <222> not found for SEQ ID#:17
L:1118 M:258 W: Mandatory Feature missing, <223> not found for SEQ ID#:17
L:1118 M:340 W: (46) "n" or "Xaa" used: Feature required, for SEQ ID#:17